

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
27 December 2001 (27.12.2001)

PCT

(10) International Publication Number  
**WO 01/97808 A1**

- (51) International Patent Classification<sup>7</sup>: **A61K 31/425**, 45/06, A61P 3/06
- (21) International Application Number: **PCT/GB01/02696**
- (22) International Filing Date: **19 June 2001 (19.06.2001)**
- (25) Filing Language: **English**
- (26) Publication Language: **English**
- (30) Priority Data:  
0014969.0 **19 June 2000 (19.06.2000)** **GB**
- (71) Applicants (*for all designated States except US*): **SMITHKLINE BEECHAM PLC [GB/GB]**; New Horizons Court, Brentford, Middlesex TW8 9EP (GB). **SMITHKLINE BEECHAM CORPORATION [US/US]**; P.O. Box 7929, Philadelphia, PA 19103 (US).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): **ARCH, Jonathan, Robert, Sanders [GB/GB]**; GlaxoSmithKline, New Frontiers Science Park South, Third Avenue, Harlow, Essex CM19 5AW (GB). **LENHARD, James, Martin [US/US]**; GlaxoSmithKline, Five Moore Drive, Research Triangle Park, NC 27709 (US).
- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
- Published:**
- *with international search report*
  - *before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments*
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*



**WO 01/97808 A1**

(54) Title: COMBINATIONS OF DEPEPTIDYL PEPTIDASE IV INHIBITORS AND OTHER ANTIDIABETIC AGENTS FOR THE TREATMENT OF DIABETE MELLITUS

(57) Abstract: A method for the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus in a mammal such as a human, which method comprises administering an effective, non-toxic and pharmaceutically acceptable amount of a dipeptidyl peptidase IV inhibitor and another antidiabetic agent, to a mammal in need thereof.

## COMBINATIONS OF DEPEPTIDYL PEPTIDASE IV INHIBITORS AND OTHER ANTIDIABETIC AGENTS FOR THE TREATMENT OF DIABETE MELLITUS

5 This invention relates to a method of treatment, in particular to a method for the treatment of diabetes mellitus, especially non-insulin dependent diabetes (NIDDM) or Type 2 diabetes and conditions associated with diabetes mellitus and to compositions for use in such method.

Dipeptidyl peptidase IV (DPP-IV) is a post-proline/alanine cleaving serine protease found in various tissues of the body including kidney, liver, and intestine.

10 It is known that DPP-IV inhibitors may be useful for the treatment of impaired glucose tolerance and diabetes mellitus (International Patent Application, Publication Number WO99/61431, Pederson RA et al, Diabetes. 1998 Aug;47(8):1253-8 and Pauly RP et al, Metabolism 1999 Mar;48(3):385-9). In particular WO99/61431 discloses DPP-IV inhibitors comprising an amino acid and a thiazolidine or pyrrolidine group, and salts thereof, such as isoleucyl (or isoleucine) thiazolidide and salts thereof.

15 Other DPP-IV inhibitors include those disclosed in United States Patent Numbers 6124305 and US 6107317, International Patent Applications, Publication Numbers WO 9819998, WO 9515309 and WO 9818763.

20 Alpha glucosidase inhibitor antihyperglycaemic agents (or alpha glucosidase inhibitors) and biguanide antihyperglycaemic agents (or biguanides) are commonly used in the treatment of Type 2 diabetes. Acarbose, voglibose, emiglitazone and miglitol are examples of alpha glucosidase inhibitors. 1,1 - Dimethylbiguanidine (or metformin) is a particular example of a biguanide.

25 Insulin secretagogues are compounds that promote increased secretion of insulin by the pancreatic beta cells. The sulphonylureas are well known examples of insulin secretagogues. The sulphonylureas act as hypoglycaemic agents and are used in the treatment of Type 2 diabetes. Examples of sulphonylureas include glibenclamide (or glyburide), glipizide, gliclazide, glimepiride, tolazamide and tolbutamide.

30 European Patent Application, Publication Number 0,306,228 relates to certain thiazolidinedione derivatives disclosed as having antihyperglycaemic and hypolipidaemic activity. One particular thiazolidinedione disclosed in EP 0306228 is 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione (hereinafter 'Compound (I)'). WO94/05659 discloses certain salts of Compound (I) including the maleate salt at example 1 thereof.

35 Compound (I) is an example of a class of anti-hyperglycaemic agents known as 'insulin sensitisers'. In particular Compound (I) is a thiazolidinedione insulin sensitiser. Compound (I) is also a peroxisome proliferator-activated receptor (PPAR $\gamma$ ) agonist insulin sensitiser.

40 European Patent Applications, Publication Numbers: 0008203, 0139421, 0032128, 0428312, 0489663, 0155845, 0257781, 0208420, 0177353, 0319189, 0332331, 0332332, 0528734, 0508740; International Patent Application, Publication Numbers 92/18501, 93/02079, 93/22445 and United States Patent Numbers 5104888 and 5478852, also disclose certain thiazolidinedione insulin sensitisers.

Another series of compounds generally recognised as having insulin sensitiser activity are those typified by the compounds disclosed in International Patent Applications, Publication Numbers WO93/21166 and WO94/01420. These compounds are herein referred to as 'acyclic insulin sensitisers'. Other examples of acyclic insulin sensitisers are those disclosed in United States Patent Number 5232945 and International Patent Applications, Publication Numbers WO92/03425 and WO91/19702.

Examples of other insulin sensitisers are those disclosed in European Patent Application, Publication Number 0533933, Japanese Patent Application Publication Number 05271204 and United States Patent Number 5264451.

10 The above mentioned publications are incorporated herein by reference.

It is now indicated that dipeptidyl peptidase IV inhibitors, such as the compounds of WO99/61431, in combination with other antidiabetic agents provide a particularly beneficial effect on glycaemic control and that such combination is therefore suggested to be particularly useful for the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus. Such combinations will provide improved blood glucose regulation without introducing unacceptable side-effects.

Accordingly, the invention provides a method for the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus in a mammal such as a human, which method comprises administering an effective, non-toxic and pharmaceutically acceptable amount of a dipeptidyl peptidase IV inhibitor and another antidiabetic agent, to a mammal in need thereof.

In another aspect the invention provides a dipeptidyl peptidase IV inhibitor and another antidiabetic agent, for use in a method for the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus.

25 The method comprises either co-administration of a dipeptidyl peptidase IV inhibitor and another antidiabetic agent or the sequential administration thereof.

Co-administration includes administration of a formulation which includes both a DPP- IV inhibitor and the other antidiabetic agent or the essentially simultaneous administration of separate formulations of each agent.

30 In another aspect the invention provides the use of a dipeptidyl peptidase IV inhibitor and another antidiabetic agent for use in the manufacture of a composition for the treatment of obesity, diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus.

Suitably, the other antidiabetic agent comprises one or more, generally one or two, of an alpha glucosidase inhibitor, a biguanide, an insulin secretagogue or an insulin sensitiser.

Suitably, the other antidiabetic agent is selected from an alpha glucosidase inhibitor, a biguanide, an insulin secretagogue or an insulin sensitiser.

A further suitable antidiabetic agent is insulin.

40 A suitable alpha glucosidase inhibitor is acarbose.

Other suitable alpha glucosidase inhibitors are emiglitate and miglitol. A further suitable alpha glucosidase inhibitor is voglibose.

Suitable biguanides include metformin, buformin or phenformin, especially metformin.

Suitable insulin secretagogues include sulphonylureas.

Suitable sulphonylureas include glibenclamide, glipizide, gliclazide, glimepiride, 5 tolazamide and tolbutamide. Further sulphonylureas include acetohexamide, carbutamide, chlorpropamide, glibornuride, gliquidone, glisentide, glisolamide, glisoxepide, glycropyamide and glycylamide. Also included is the sulphonylurea glipentide.

A further suitable insulin secretagogue is repaglinide. An additional insulin 10 secretagogue is nateglinide.

Insulin sensitisers include PPAR $\gamma$  agonist insulin sensitisers including the compounds disclosed in WO 97/31907 and especially 2-(1-carboxy-2-{4-{2-(5-methyl-2-phenyl-oxazol-4-yl)-ethoxy}-phenyl}-ethylamino)-benzoic acid methyl ester and 2(S)-(2-benzoyl-phenylamino)-3-{4-[2-(5-methyl-2-phenyl-oxazol-4-yl)-ethoxy]-phenyl}- 15 propionic acid.

Insulin sensitisers also include thiazolidinedione insulin sensitisers.

A preferred insulin sensitiser is Compound (I) or a derivative thereof.

Other suitable thiazolidinedione insulin sensitisers include (+) -5-[[4-[(3,4-dihydro-6-hydroxy-2,5,7,8-tetramethyl-2H-1-benzopyran-2-yl)methoxy]phenyl]methyl]- 20 2,4-thiazolidinedione (or troglitazone), 5-[4-[(1-methylcyclohexyl)methoxy]benzyl]thiazolidine-2,4-dione (or ciglitazone), 5-[4-[2-(5-ethylpyridin-2-yl)ethoxy]benzyl]thiazolidine-2,4-dione (or pioglitazone) or 5-[(2-benzyl-2,3-dihydrobenzopyran)-5-ylmethyl]thiazolidine-2,4-dione (or englitazone).

A particular thiazolidinedione insulin sensitiser is 5-[4-[2-(5-ethylpyridin-2-yl)ethoxy]benzyl] thiazolidine-2,4-dione (or pioglitazone). 25

A particular thiazolidinedione insulin sensitiser is (+) -5-[[4-[(3,4-dihydro-6-hydroxy-2,5,7,8-tetramethyl-2H-1-benzopyran-2-yl)methoxy]phenyl]methyl]-2,4-thiazolidinedione (or troglitazone).

Particular DPP- IV inhibitors include the specific examples disclosed in 30 WO99/61431, such as L-threo-isoleucyl pyrrolidide, L-allo-isoleucyl thiazolidide, L-allo-isoleucyl pyrrolidide and salts thereof. A particular DPP- IV inhibitor is isoleucine thiazolidide and salts thereof.

Further DPP-IV inhibitors include the specific examples disclosed in United States Patent Numbers 6124305 and US 6107317, International Patent Applications, 35 Publication Numbers WO 9819998, WO 9515309 and WO 9818763; such as 1[2-[(5-cyanopyridin-2-yl)aminoethylamino]acetyl-2-cyano-(S)-pyrrolidine and (2S)-1-[(2S)-2-amino-3,3-dimethylbutanoyl]-2-pyrrolidinecarbonitrile.

For the avoidance of doubt, the examples disclosed in each of the above mentioned publications are specifically incorporated herein by reference, as individually 40 disclosed compounds.

It will be understood that the DPP- IV inhibitor and the other antidiabetic agent are each administered in a pharmaceutically acceptable form, including pharmaceutically acceptable derivatives such as pharmaceutically acceptable salts, esters and solvates

thereof, as appropriate of the relevant pharmaceutically active agent. In certain instances herein the names used for the other antidiabetic agent may relate to a particular pharmaceutical form of the relevant active agent: It will be understood that all pharmaceutically acceptable forms of the active agents per se are encompassed by this invention.

Suitable pharmaceutically acceptable forms of the other antidiabetic agent depend upon the particular agent being used but include known pharmaceutically acceptable forms of the particular agent chosen. Such derivatives are found or are referred to in standard reference texts such as the British and US Pharmacopoeias, Remington's Pharmaceutical Sciences (Mack Publishing Co.), Martindale The Extra Pharmacopoeia (London, The Pharmaceutical Press) (for example see the 31st Edition page 341 and pages cited therein) or the above mentioned publications.

Suitable pharmaceutically acceptable forms of the DPP- IV inhibitor include salted forms and solvated forms, include those described in WO 99/61431, for example the fumarate salt

The DPP- IV inhibitor is prepared according to published methods, for example when the DPP- IV inhibitor is a compound of WO 99/61431 or a derivative thereof such as a pharmaceutically acceptable salt thereof or a pharmaceutically acceptable solvate thereof, then it is prepared according to methods disclosed therein. Similarly for the compounds of United States Patent Numbers 6124305 and US 6107317 and those of International Patent Applications, Publication Numbers WO 9819998, WO 9515309 and WO 9818763.

Certain of the compounds mentioned herein may contain one or more chiral carbon atoms and hence can exist in two or more isomeric forms, all of which are encompassed by the invention, either as individual isomers or as mixtures of isomers, including racemates. Certain of the compounds mentioned herein, in particular the thiazolidinediones such as Compound (I), may exist in one of several tautomeric forms, all of which are encompassed by the invention as individual tautomeric forms or as mixtures thereof

The DPP- IV inhibitor and the other antidiabetic agent of choice is prepared according to known methods, such methods are found or are referred to in standard reference texts, such as the British and US Pharmacopoeias, Remington's Pharmaceutical Sciences (Mack Publishing Co.), Martindale The Extra Pharmacopoeia (London, The Pharmaceutical Press) (for example see the 31st Edition page 341 and pages cited therein) or the above mentioned publications.

When used herein the term 'conditions associated with diabetes' includes those conditions associated with the pre-diabetic state, conditions associated with diabetes mellitus itself and complications associated with diabetes mellitus.

When used herein the term 'conditions associated with the pre-diabetic state' includes conditions such as insulin resistance, including hereditary insulin resistance, impaired glucose tolerance and hyperinsulinaemia.

'Conditions associated with diabetes mellitus itself' include hyperglycaemia, insulin resistance, including acquired insulin resistance and obesity. Further conditions

associated with diabetes mellitus itself include hypertension and cardiovascular disease, especially atherosclerosis and conditions associated with insulin resistance. Conditions associated with insulin resistance include polycystic ovarian syndrome and steroid induced insulin resistance and gestational diabetes.

- 5        'Complications associated with diabetes mellitus' includes renal disease, especially renal disease associated with Type 2 diabetes, neuropathy and retinopathy.

Renal diseases associated with Type 2 diabetes include nephropathy, glomerulonephritis, glomerular sclerosis, nephrotic syndrome, hypertensive nephrosclerosis and end stage renal disease.

- 10       As used herein the term 'pharmaceutically acceptable' embraces both human and veterinary use: for example the term 'pharmaceutically acceptable' embraces a veterinarily acceptable compound.

Diabetes mellitus is preferably Type 2 diabetes.

- 15       Suitably, the particularly beneficial effect on glycaemic control provided by the treatment of the invention is an improved therapeutic ratio for the combination of the invention relative to the therapeutic ratio for one compound of the combination when used alone and at a dose providing an equivalent efficacy to the combination of the invention.

- 20       In a preferred aspect, the particularly beneficial effect on glycaemic control provided by the treatment of the invention is indicated to be a synergistic effect relative to the control expected from the effects of the individual active agents.

In a further aspect of the invention, combining doses of the DPP- IV inhibitor and the other agent will produce a greater beneficial effect than can be achieved for either agent alone at a dose twice that used for that agent in the combination.

- 25       Glycaemic control may be characterised using conventional methods, for example by measurement of a typically used index of glycaemic control such as fasting plasma glucose or glycosylated haemoglobin (Hb A1c). Such indices are determined using standard methodology, for example those described in: Tuescher A, Richterich, P., Schweiz. med. Wschr. 101 (1971), 345 and 390 and Frank P., 'Monitoring the Diabetic  
30       Patent with Glycosolated Hemoglobin Measurements', Clinical Products 1988.

In a preferred aspect, the dosage level of each of the active agents when used in accordance with the treatment of the invention will be less than would have been required from a purely additive effect upon glycaemic control.

- 35       It is also considered that the treatment of the invention will effect an improvement, relative to the individual agents, in the levels of advanced glycosylation end products (AGEs), and serum lipids including total cholesterol, HDL-cholesterol, LDL-cholesterol including improvements in the ratios thereof, in particular an improvement in serum lipids including total cholesterol, HDL-cholesterol, LDL-cholesterol including improvements in the ratios thereof.

- 40       In the treatment of the invention, the active medicaments are preferably administered in pharmaceutical composition form. As indicated above, such compositions can include both medicaments or one only of the medicaments.

Accordingly, in one aspect the present invention also provides a pharmaceutical composition comprising a dipeptidyl peptidase IV inhibitor and another antidiabetic agent and a pharmaceutically acceptable carrier therefor.

Thus, in a further aspect, the invention also provides a process for preparing a  
5 pharmaceutical composition comprising a dipeptidyl peptidase IV inhibitor, another antidiabetic agent and a pharmaceutically acceptable carrier therefor, which process comprises admixing the dipeptidyl peptidase IV inhibitor, another antidiabetic agent and a pharmaceutically acceptable carrier.

The compositions are preferably in a unit dosage form in an amount appropriate  
10 for the relevant daily dosage.

Suitable dosages, including especially unit dosages, of the DPP- IV inhibitor or the other antidiabetic agent include the known dosages including unit doses for these compounds as described or referred to in reference text such as the British and US Pharmacopoeias, Remington's Pharmaceutical Sciences (Mack Publishing Co.),  
15 Martindale The Extra Pharmacopoeia (London, The Pharmaceutical Press) (for example see the 31st Edition page 341 and pages cited therein) or the above mentioned publications.

Thus, suitable dosages for the DPP- IV inhibitors of WO 99/61431 and include those disclosed therein, for example 0.01 to 30mg per day or 0.01 to 10mg per kilogram  
20 of body weight. Also, the suitable doses of the other DPP- IV inhibitors mentioned herein include those mentioned in the relevant publications mentioned above.

For the alpha glucosidase inhibitor, a suitable amount of acarbose is in the range of from 25 to 600 mg, including 50 to 600 mg, for example 100mg or 200mg.

For the biguanide, a suitable dosage of metformin is between 100 to 3000mg, for  
25 example 250, 500mg, 850mg or 1000mg.

For the insulin secretagogue, a suitable amount of glibenclamide is in the range of from 2.5 to 20 mg, for example 10mg or 20mg; a suitable amount of glipizide is in the range of from 2.5 to 40 mg; a suitable amount of gliclazide is in the range of from 40 to 320 mg; a suitable amount of tolazamide is in the range of from 100 to 1000 mg; a  
30 suitable amount of tolbutamide is in the range of from 1000 to 3000 mg; a suitable amount of chlorpropamide is in the range of from 100 to 500 mg; and a suitable amount of gliquidone is in the range of from 15 to 180 mg. Also a suitable amount of glimepiride is 1 to 6mg and a suitable amount of glipentide is 2.5 to 20mg.

A suitable amount of repaglinide is in the range of from 0.5mg to 20mg, for  
35 example 16mg. Also a suitable amount of nateglinide is 90 to 360mg, for example 270mg.

In one particular aspect, the composition comprises 2 to 12 mg of Compound (I).

Suitably the composition comprises 2, 3, 4, 5, 6, 7, 8, 9, 10, 11 or 12 mg of Compound (I).

Particularly, the composition comprises 2 to 4 , 4 to 8 or 8 to 12 mg of  
40 Compound (I).

Particularly, the composition comprises 2 to 4mg of Compound (I).

Particularly, the composition comprises 4 to 8mg of Compound (I).

Particularly, the composition comprises 8 to 12 mg of Compound (I).

Preferably, the composition comprises 2 mg of Compound (I).

Preferably, the composition comprises 4 mg of Compound (I).

Preferably, the composition comprises 8 mg of Compound (I).

5        Suitable unit dosages of other insulin sensitisers include from 100 to 800mg of troglitazone such as 200, 400, 600 or 800mg or from 5 to 50mg, including 10 to 40mg, of pioglitazone, such as 20, 30 or 40 mg and also including 15, 30 and 45mg of pioglitazone.

      Suitable dosages of other PPAR $\gamma$  agonist insulin sensitisers include those disclosed for the respective agonist in the abovementioned applications, for example 2-(1-  
10        carboxy-2-{4-[2-(5-methyl-2-phenyl-oxazol-4-yl)-ethoxy]-phenyl}-ethylamino)-benzoic acid methyl ester and 2(S)-(2-benzoyl-phenylamino)-3-{4-[2-(5-methyl-2-phenyl-oxazol-4-yl)-ethoxy]-phenyl}-propionic acid are suitably dosed in accordance with the dosages disclosed in WO 97/31907.

      In the treatment the medicaments may be administered from 1 to 6 times a day,  
15        but most preferably 1 or 2 times per day.

      Also, the dosages of each particular active agent in any given composition can as required vary within a range of doses known to be required in respect of accepted dosage regimens for that compound. Dosages of each active agent can also be adapted as required to take into account advantageous effects of combining the agents as mentioned  
20        herein.

      It will be understood that the DPP- IV inhibitor and the other antidiabetic agent are in a pharmaceutically acceptable form, including pharmaceutically acceptable derivatives such as pharmaceutically acceptable salts, esters and solvates thereof, as appropriate to the relevant pharmaceutically active agent chosen. In certain instances  
25        herein the names used for the antidiabetic agent may relate to a particular pharmaceutical form of the relevant active agent: It will be understood that all pharmaceutically acceptable forms of the active agents per se are encompassed by this invention.

      The present invention also provides a pharmaceutical composition comprising a dipeptidyl peptidase IV inhibitor, another antidiabetic agent and a pharmaceutically  
30        acceptable carrier therefor, for use as an active therapeutic substance.

      In particular, the present invention provides a pharmaceutical composition comprising a dipeptidyl peptidase IV inhibitor, another antidiabetic agent and a pharmaceutically acceptable carrier therefor, for use in the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus.

35        Usually the compositions are adapted for oral administration. However, they may be adapted for other modes of administration, for example parenteral administration, sublingual or transdermal administration.

      The compositions may be in the form of tablets, capsules, powders, granules, lozenges, suppositories, reconstitutable powders, or liquid preparations, such as oral or  
40        sterile parenteral solutions or suspensions.

      In order to obtain consistency of administration it is preferred that a composition of the invention is in the form of a unit dose.

Unit dosage presentation forms for oral administration may be in tablet or capsule form and may as necessary contain conventional excipients such as binding agents, fillers, lubricants, glidants, disintegrants and wetting agents.

5 The solid oral compositions may be prepared by conventional methods of blending, filling or tableting. Repeated blending operations may be used to distribute the active agent throughout those compositions employing large quantities of fillers. Such operations are of course conventional in the art. The tablets may be coated according to methods well known in normal pharmaceutical practice, in particular with an enteric coating.

10 Oral liquid preparations may be in the form of, for example, emulsions, syrups, or elixirs, or may be presented as a dry product for reconstitution with water or other suitable vehicle before use. Such liquid preparations may contain conventional additives such as suspending agents, for example sorbitol, syrup, methyl cellulose, gelatin, hydroxyethylcellulose, carboxymethylcellulose, aluminium stearate gel, hydrogenated  
15 edible fats; emulsifying agents, for example lecithin, sorbitan monooleate, or acacia; non-aqueous vehicles (which may include edible oils), for example almond oil, fractionated coconut oil, oily esters such as esters of glycerine, propylene glycol, or ethyl alcohol; preservatives, for example methyl or propyl p-hydroxybenzoate or sorbic acid; and if desired conventional flavouring or colouring agents.

20 For parenteral administration, fluid unit dosage forms are prepared utilizing the compound and a sterile vehicle, and, depending on the concentration used, can be either suspended or dissolved in the vehicle. In preparing solutions the compound can be dissolved in water for injection and filter sterilized before filling into a suitable vial or ampoule and sealing. Advantageously, adjuvants such as a local anaesthetic, a  
25 preservative and buffering agent can be dissolved in the vehicle. To enhance the stability, the composition can be frozen after filling into the vial and the water removed under vacuum. Parenteral suspensions are prepared in substantially the same manner, except that the active compound is suspended in the vehicle instead of being dissolved, and sterilization cannot be accomplished by filtration. The compound can be sterilized by  
30 exposure to ethylene oxide before suspending in the sterile vehicle. Advantageously, a surfactant or wetting agent is included in the composition to facilitate uniform distribution of the compound.

Compositions may contain from 0.1% to 99% by weight, preferably from 10-60% by weight, of the active material, depending upon the method of administration.

35 Examples of binding agents include acacia, alginic acid, carboxymethylcellulose calcium, carboxymethylcellulose sodium, dextrates, dextrin, dextrose, ethylcellulose, gelatin, liquid glucose, guar gum, hydroxyethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methylcellulose, magnesium aluminium silicate, maltodextrin, methyl cellulose, polymethacrylates, polyvinylpyrrolidone, pregelatinised starch, sodium  
40 alginate, sorbitol, starch, syrup, tragacanth.

Examples of fillers include calcium carbonate, calcium phosphate, calcium sulphate, carboxymethylcellulose calcium, carboxymethylcellulose sodium, compressible sugar, confectioner's sugar, dextrates, dextrin, dextrose, dibasic calcium phosphate

5 dihydrate, dibasic calcium phosphate, fructose, glyceryl palmitostearate, glycine, hydrogenated vegetable oil-type 1, kaolin, lactose, maize starch, magnesium carbonate, magnesium oxide, maltodextrin, mannitol, microcrystalline cellulose, polymethacrylates, potassium chloride, powdered cellulose, pregelatinised starch, sodium chloride, sorbitol, starch, sucrose, sugar spheres, talc, tribasic calcium phosphate, xylitol.

Examples of lubricants include calcium stearate, glyceryl monostearate, glyceryl palmitostearate, magnesium stearate, microcrystalline cellulose, sodium benzoate, sodium chloride, sodium lauryl sulphate, stearic acid, sodium stearyl fumarate, talc, zinc stearate.

10 Examples of glidants include colloidal silicon dioxide, powdered cellulose, magnesium trisilicate, silicon dioxide, talc.

Examples of disintegrants include alginic acid, carboxymethylcellulose calcium, carboxymethylcellulose sodium, colloidal silicon dioxide, croscarmellose sodium, crospovidone, guar gum, magnesium aluminium silicate, microcrystalline cellulose, methyl cellulose, polyvinylpyrrolidone, polacrillin potassium, pregelatinised starch, sodium alginate, sodium lauryl sulphate, sodium starch glycollate.

15 An example of a pharmaceutically acceptable wetting agent is sodium lauryl sulphate.

The compositions are prepared and formulated according to conventional methods, such as those disclosed in standard reference texts, for example the British and US Pharmacopoeias, Remington's Pharmaceutical Sciences (Mack Publishing Co.), Martindale The Extra Pharmacopoeia (London, The Pharmaceutical Press) (for example see the 31st Edition page 341 and pages cited therein) and Harry's Cosmeticology (Leonard Hill Books) or the above mentioned publications.

25 For example, the solid oral compositions may be prepared by conventional methods of blending, filling or tableting. Repeated blending operations may be used to distribute the active agent throughout those compositions employing large quantities of fillers. Such operations are of course conventional in the art. The tablets may be coated according to methods well known in normal pharmaceutical practice.

30 Compositions may, if desired, be in the form of a pack accompanied by written or printed instructions for use.

No adverse toxicological effects are expected for the compositions or methods of the invention in the above mentioned dosage ranges.

### Pharmacological Data

Age and weight matched male ZDF *fa/fa* rats (Genetic Models, Inc., Indianapolis, IN) were housed individually at 72° F and 50% relative humidity with a 12 h light/dark cycle and fed PMI 5008 Formulab Diet (PMI Nutrition International, Saint Louis, MO).

- 5        Animals were dosed by oral gavage twice daily during the dark cycle for one week with vehicle (0.5% hydroxy-propylmethylcellulose (HPMC) plus 0.1% Tween 80), 100 mg/kg isoleucine thiazolidide (Compound (II)), 5 mg/kg Compound (I) in vehicle, or 5 mg/kg Compound (I) plus 100 mg/kg Compound (II) in vehicle.

- 10       For glucose tolerance measurements, rats were treated with test compound for 7 days and given an intraperitoneal injection of a glucose solution in saline 30 minutes after the last dose of test compound.

- 15       Rats were anesthetized with isoflurane for cardiac blood collection 30 minutes after administration of the glucose solution. Serum chemistry measurements were obtained using an automated chemistry analyzer (ILab600, Instrument Laboratory, Lexington, MA).

- 20       DPP-IV activity was measured using the fluorogenic substrate Gly-Pro-AMC (50 mM) according to the manufacturers specification (Enzyme System Products, Livermore CA). The substrate was mixed with 50 mM Tris, pH 7.8, in plasma (20% final v/v) and the samples were incubated for 5-20 min at 30°C. DPP-IV activity was determined by measuring fluorescence using a cytofluor spectrofluorometer with the filters set at 360 nm excitation and 460 nm emission.

Results from each group (n=6) were averaged and compared to vehicle treated rats to determine significance and are shown in Table I.

The following data illustrates the invention but does not limit it in any way.

TABLE I

5 ZDF rats, treated BID for 7 days

	Plasma DPP-IV activity	% HbA1C (7 day change)	Plasma Glucose (30 min GTT)
Control	5544 ± 485	1.63 ± 1.12	695 ± 24
Compound (I) (5 mg/kg)	4104 ± 399*	0.79 ± 0.54*	665 ± 40
Compound (II) (100 mg/kg)	962 ± 53*	1.81 ± 1.24	684 ± 60
Combination	703 ± 16*	0.41 ± 0.28*	454 ± 52*

\*P.0.05

## CLAIMS

1. A method for the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus in a mammal such as a human, which method comprises administering an effective, non-toxic and pharmaceutically acceptable amount of a dipeptidyl peptidase IV inhibitor and another antidiabetic agent, to a mammal in need thereof.
2. A method according to claim 1, wherein the dipeptidyl peptidase IV is selected from: L-threo-isoleucyl pyrrolidide, L-allo-isoleucyl thiazolidide, L-allo-isoleucyl pyrrolidide; and salts thereof or 1[2-[(5-cyanopyridin-2-yl)aminoethylamino]acetyl-2-cyano-(S)-pyrrolidine and (2S)-1-[(2S)-2-amino-3,3-dimethylbutanoyl]-2-pyrrolidinecarbonitrile.
3. A method according to claim 1 or claim 2, wherein the other antidiabetic agent comprises one or more of an alpha glucosidase inhibitor, a biguanide, an insulin secretagogue or an insulin sensitiser.
3. A method according to claim 3, wherein the alpha glucosidase inhibitor is selected from acarbose, emiglitate, miglitol and voglibose.
4. A method according to claim 3, wherein the biguanide is selected from metformin, buformin and phenformin.
5. A method according to claim 3, wherein the insulin secretagogue is a sulphonylurea selected from: glibenclamide, glipizide, gliclazide, glimepiride, tolazamide and tolbutamide, acetohexamide, carbutamide, chlorpropamide, glibornuride, gliquidone, glisentide, glisolamide, glisoxepide, glycopyramide, glycylamide and glipentide.
6. A method according to claim 3, wherein the insulin secretagogue is repaglinide or nateglinide.
7. A method according to claim 3, wherein the insulin sensitiser is thiazolidinedione.
8. A method according to claim 7, wherein the thiazolidinedione is selected from: (+) -5-[[4-[(3,4-dihydro-6-hydroxy-2, 5, 7, 8-tetramethyl-2H-1-benzopyran-2-yl)methoxy]phenyl]methyl]-2,4-thiazolidinedione (or troglitazone), 5-[4-[(1-methylcyclohexyl)methoxy]benzyl] thiazolidine-2,4-dione (or ciglitazone), 5-[4-[2-(5-ethylpyridin-2-yl)ethoxy]benzyl] thiazolidine-2,4-dione (or pioglitazone) or 5-[(2-benzyl-2,3-dihydrobenzopyran)-5-ylmethyl]thiazolidine-2,4-dione (or englitazone).

9. A method according to claim 7, wherein the thiazolidinedione is 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione, 5-[4-[2-(5-ethylpyridin-2-yl)ethoxy]benzyl] thiazolidine-2,4-dione (or pioglitazone); or a pharmaceutically acceptable derivative thereof.
- 5
10. A pharmaceutical composition comprising a dipeptidyl peptidase IV inhibitor and another antidiabetic agent and a pharmaceutically acceptable carrier therefor.
11. A process for preparing a pharmaceutical composition comprising a dipeptidyl peptidase IV inhibitor, another antidiabetic agent and a pharmaceutically acceptable carrier therefor, which process comprises admixing the dipeptidyl peptidase IV inhibitor, the other antidiabetic agent and a pharmaceutically acceptable carrier.
- 10
12. A pharmaceutical composition comprising a dipeptidyl peptidase IV inhibitor, another antidiabetic agent and a pharmaceutically acceptable carrier therefor, for use as an active therapeutic substance.
- 15
13. A pharmaceutical composition comprising a dipeptidyl peptidase IV inhibitor, another antidiabetic agent and a pharmaceutically acceptable carrier therefor, for use in the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus.
- 20

## INTERNATIONAL SEARCH REPORT

International Application No

PC 01/02696

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K31/425 A61K45/06 A61P3/06

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data, EPO-Internal

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E	WO 01 52825 A (NOVARTIS ERFINDE VERWALT GMBH ;NOVARTIS AG (CH); HOLMES DAVID GRENV) 26 July 2001 (2001-07-26) the whole document	1-13
X	WO 99 61431 A (GLUND KONRAD ;KRUBER SUSANNE (DE); DEMUTH HANS ULRICH (DE); PROBIO) 2 December 1999 (1999-12-02) cited in the application	1-13
Y	page 8, line 13-15 claim 14	1-3, 9-13



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

## \* Special categories of cited documents:

\*A\* document defining the general state of the art which is not considered to be of particular relevance

\*E\* earlier document but published on or after the international filing date

\*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

\*O\* document referring to an oral disclosure, use, exhibition or other means

\*P\* document published prior to the international filing date but later than the priority date claimed

\*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

\*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*G\* document member of the same patent family

Date of the actual completion of the international search

12 October 2001

Date of mailing of the international search report

25/10/2001

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax (+31-70) 340-3016

Authorized officer

Veronese, A

## INTERNATIONAL SEARCH REPORT

International Application No

PC1768 01/02696

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>PAULY R P ET AL: "IMPROVED GLUCOSE TOLERANCE IN RATS TREATED WITH THE DIPEPTIDYL PEPTIDASE IV (CD26) INHIBITOR ILE-THIAZOLIDIDE" METABOLISM, CLINICAL AND EXPERIMENTAL, W.B. SAUNDERS CO., PHILADELPHIA, PA, US, vol. 48, no. 3, 1999, pages 385-389, XP000874051 ISSN: 0026-0495 See results and discussion figure 2</p>	1-3, 10-13
X	<p>DEACON C F ET AL: "DIPEPTIDYL PEPTIDASE IV INHIBITION POTENTIATES THE INSULINOTROPIC EFFECT OF GLUCAGON-LIKE PEPTIDE 1 IN THE ANESTHETIZED PIG" DIABETES, NEW YORK, NY, US, vol. 47, no. 5, May 1998 (1998-05), pages 764-769, XP000853618 ISSN: 0012-1797 figures; tables</p>	1,10-13
X	<p>HOLST J J ET AL: "INHIBITION OF THE ACTIVITY OF DIPEPTIDYL-PEPTIDASE IV AS A TREATMENT FOR TYPE 2 DIABETES" DIABETES, NEW YORK, NY, US, vol. 47, November 1998 (1998-11), pages 1663-1670, XP000853619 ISSN: 0012-1797 See discussion and conclusions page 2</p>	1,10-13
Y	<p>EP 0 306 228 A (BEECHAM GROUP PLC) 8 March 1989 (1989-03-08) claims 16,17; example 5</p>	1-13
Y	<p>WO 98 19998 A (CIBA GEIGY AG ;VILLHAUER EDWIN BERNARD (US)) 14 May 1998 (1998-05-14) page 18, line 17-23; claims; example 3</p>	1-13

## FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 3,4,5,6

Present claims 1-13 relate to a very large number of possible compounds and compositions. Support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT is to be found, however, for only a very small proportion of the compounds and the compositions claimed. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Consequently, the search has been carried out for those parts of the claims which appear to be supported and disclosed, namely the example provided at page 10 and 11, disclosing compositions comprising the association of compound I and compound II, and the general idea underlying the application (the idea to associate a dipeptidyl peptidase IV inhibitor (DPP IV) and an other antidiabetic agent). In view of the prior art retrieved during the search, disclosing the use of DPP IV inhibitors together with other antidiabetic agents, the application would seem not to be linked by a common inventive concept and would not meet the requirements of unity (Art. 3.4(iii) and Rule 13 PCT). Different inventions can be defined, each of them identified by the combination of the different DPP IV inhibitors with the different classes of antidiabetic agents. As already mentioned above, in the present application only a small proportion of the claimed subject matter is disclosed and supported, and therefore only the invention for which support has been provided has been searched. A non-unity objection has not been raised for reasons of procedural economy.

An error in the claim numbering is apparent. Two claims 3 are present. Only the first one has been searched (incompletely).

Claims searched incompletely: 1-3, 7-13

Not searched claims: 3-6

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

## INTERNATIONAL SEARCH REPORT

Int — nal Application No

PCT/GB 01/02696

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 0152825	A	26-07-2001	WO 0152825 A2	26-07-2001
WO 9961431	A	02-12-1999	DE 19823831 A1	02-12-1999
			AU 4370999 A	13-12-1999
			BR 9910758 A	13-02-2001
			CN 1303381 T	11-07-2001
			DE 29909208 U1	09-09-1999
			DE 29909210 U1	09-09-1999
			DE 29909211 U1	23-09-1999
			WO 9961431 A1	02-12-1999
			EP 1082314 A1	14-03-2001
			NO 20005994 A	25-01-2001
EP 0306228	A	08-03-1989	AT 186724 T	15-12-1999
			AU 2173888 A	09-03-1989
			BR 1100841 A3	20-06-2000
			CA 1328452 A1	12-04-1994
			CA 1339902 A1	09-06-1998
			CZ 9103916 A3	17-03-1993
			DE 3856378 D1	23-12-1999
			DE 3856378 T2	11-05-2000
			DK 490288 A	05-03-1989
			DK 200001556 A	18-10-2000
			EP 0306228 A1	08-03-1989
			EP 0842925 A1	20-05-1998
			ES 2137915 T3	01-01-2000
			GR 3031873 T3	29-02-2000
			HK 1011029 A1	03-11-2000
			JP 10194970 A	28-07-1998
			JP 10194971 A	28-07-1998
			JP 1131169 A	24-05-1989
			JP 2614497 B2	28-05-1997
			JP 2817840 B2	30-10-1998
			JP 9183771 A	15-07-1997
			JP 2837139 B2	14-12-1998
			JP 9183726 A	15-07-1997
			JP 9183772 A	15-07-1997
			KR 164207 B1	15-01-1999
			KR 164275 B1	15-01-1999
			KR 169463 B1	15-01-1999
			LU 90711 A9	05-03-2001
			NZ 226027 A	26-03-1992
			PT 88410 A ,B	31-07-1989
			SG 59988 A1	22-02-1999
			SK 391691 A3	11-12-2000
			US 6288095 B1	11-09-2001
			US 5646169 A	08-07-1997
			US 5002953 A	26-03-1991
			US 5521201 A	28-05-1996
			US 5232925 A	03-08-1993
			US 5194443 A	16-03-1993
			US 5756525 A	26-05-1998
			US 5260445 A	09-11-1993
			ZA 8806536 A	26-07-1989
WO 9819998	A	14-05-1998	AU 726186 B2	02-11-2000
			AU 5318498 A	29-05-1998
			BR 9714130 A	29-02-2000

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 01/02696

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9819998	A	CN 1236361 A	24-11-1999
		CZ 9901615 A3	11-08-1999
		WO 9819998 A2	14-05-1998
		EP 0937040 A2	25-08-1999
		HU 0000323 A2	28-08-2000
		JP 2000511559 T	05-09-2000
		NO 992028 A	28-04-1999
		PL 332777 A1	11-10-1999
		SK 60899 A3	10-04-2000
		TR 9901004 T2	21-07-1999